

wherein the configuration at C-2 is (R) or (S) or a mixture thereof, and
wherein R is NR₁R₂-C₂₋₆alkylene or guanidine-C₂₋₆alkylene, and each of R₁ and R₂
independently is H or C₁₋₄alkyl,
in free form, salt form or protected form.

Claim 3 (currently amended): Microparticles according to claim 1 ~~or 2~~ wherein the
somatostatin analogue is in pamoate salt form.

Claim 4 (currently amended): Microparticles according to ~~any preceding~~ claim 1
wherein the polymer matrix comprises a linear or branched polylactide-co-glycolide.

Claim 5 (currently amended): Microparticles according to ~~any preceding~~ claim 1
wherein the polymeric matrix comprises at least two different polymers.

Claim 6 (currently amended): Microparticles according to ~~any preceding~~ claim 1
further comprising a surfactant, a porosity influencing agent and/or a basic salt.

Claim 7 (currently amended): A pharmaceutical composition comprising
microparticles of ~~any preceding~~ claim 1 and a water-based vehicle comprising a wetting
agent.

Claim 8 (original): A composition according to claim 7 wherein the wetting agent
comprises a poloxamer and/or a polyoxyethylene-sorbitan-fatty acid ester.

Claim 9 (currently amended): A composition according to ~~any one of claims 7 or~~
8 claim 7 wherein the vehicle comprises a tonicity agent.

Claim 10 (currently amended): A composition according to ~~any one of claims 7 or~~
8 claim 7 wherein the vehicle comprises a viscosity increasing agent.

Claim 11 (currently amended): A kit comprising microparticles according to ~~any one~~
~~of claims 1 to 6~~ claim 1 and a water-based vehicle.

Claim 12 Canceled.

Claim 13 (currently amended): A method of treating a disease or disorder with an
aetiology comprising or associated with excess GH- and/or IGF-1 secretion in a subject in
need thereof which comprises administering microparticles according to claim 1 ~~any one of~~
~~claims 1 to 6 or of a pharmaceutical composition according to any one of claims 7 to 10 to~~
~~the subject.~~